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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/037,415	01/04/2002	David Baltimore	75723-ZB/JPW/GJG	7747
23432	7590	02/19/2010	EXAMINER	
COOPER & DUNHAM, LLP 30 Rockefeller Plaza 20th Floor NEW YORK, NY 10112			HIBBERT, CATHERINE S	
ART UNIT		PAPER NUMBER		
1636				
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary	Application No.	Applicant(s)
	10/037,415	BALTIMORE ET AL.
	Examiner	Art Unit
	CATHERINE HIBBERT	1636

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

1) Responsive to communication(s) filed on 09 November 2009.

2a) This action is **FINAL**. 2b) This action is non-final.

3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

4) Claim(s) 89 is/are pending in the application.

4a) Of the above claim(s) _____ is/are withdrawn from consideration.

5) Claim(s) _____ is/are allowed.

6) Claim(s) 89 is/are rejected.

7) Claim(s) _____ is/are objected to.

8) Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

9) The specification is objected to by the Examiner.

10) The drawing(s) filed on _____ is/are: a) accepted or b) objected to by the Examiner.

Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).

Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).

11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).

a) All b) Some * c) None of:

1. Certified copies of the priority documents have been received.
2. Certified copies of the priority documents have been received in Application No. _____.
3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

1) Notice of References Cited (PTO-892)

2) Notice of Draftsperson's Patent Drawing Review (PTO-948)

3) Information Disclosure Statement(s) (PTO/SB/08)
Paper No(s)/Mail Date 5/11/09;9/24/09;11/9/09;1/26/2010.

4) Interview Summary (PTO-413)
Paper No(s)/Mail Date. _____.

5) Notice of Informal Patent Application

6) Other: _____.

DETAILED ACTION

Applicants Amendment to the Claims filed 9 November 2009 has been received and entered. Claims 1-88 are cancelled. Claim 89 is pending and under examination.

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Any rejections not repeated in this action are WITHDRAWN.

Information Disclosure Statement

The Information Disclosure Statements filed 5/11/2009, 8/11/2008, 9/24/2009, 11/9/2009, and 1/26/2010 have been considered.

Priority

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Priority for the subject matter of claim 89 is granted back to the filing date of the 07/341,436 patent (4/21/1989).

Response to Amendment/Argument***Obviousness Type Double Patenting Rejections-maintained***

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the “right to exclude” granted by a patent and to prevent possible harassment by multiple assignees. A nonstatutory obviousness-type double patenting rejection is appropriate where the conflicting claims are not identical, but at least one examined application claim is not patentably distinct from the reference claim(s) because the examined application claim is either anticipated by, or would have been obvious over, the reference claim(s). See, e.g., *In re Berg*, 140 F.3d 1428, 46 USPQ2d 1226 (Fed. Cir. 1998); *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); and *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) or 1.321(d) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent either is shown to

be commonly owned with this application, or claims an invention made as a result of activities undertaken within the scope of a joint research agreement.

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

Claim 89 stands rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1-5, 7-17, 20-63, 88-176 and 192-203 of U.S. Patent No. 6,410,516 (hereafter the '516 patent). Although the conflicting claims are not identical, they are not patentably distinct from each other because both sets of claims read on methods of regulating NF- κ B mediated gene expression in a cell, comprising altering (inhibiting) NF- κ B activity in the cell. The instant claim is generic to the claims recited in the '516 patent. That is, the recited claims of the '516 patent fall entirely within the scope of the instant claims, or in other words, the instant claim is anticipated by the claims of the '516 patent. For example, the various methods for inhibiting expression of NF- κ B mediated gene expression in cells recited in the '516 patent are encompassed within the instant broad methodologies (i.e. the instant methods encompass any method of regulating NF- κ B mediated gene expression in any cell). With regard to instant claim 89, this claim differs from the claims in the '516 patent in reciting that the external influence is an extracellular polypeptide whereas any external influence is recited in the '516 patent claims. Since the specification of the '516 patent specifically recites extracellular polypeptides as an external influence which can induce expression of genes by inducing NF- κ B activation, it must be considered that this would have been an obvious species of external influence.

Applicants have responded to this rejection (see Applicants Remarks, filed 9 November 2009) by indicating that they will file a Terminal Disclaimer upon indication of allowable subject matter should the allowable subject matter so require.

The rejection is therefore MAINTAINED.

Claim 89 is provisionally rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 90-91 of copending Application No. 10/037,341 (hereafter the '341 application). It is noted that the '341 Application cancelled some conflicting claims and added new conflicting Claim 91 and thus this rejection has been adjusted accordingly. Although the conflicting claims are not identical, they are not patentably distinct from each other because both sets of claims recite the same methods of reducing expression in cells of a gene whose expression is modulated by NF- κ B. The instant claims differ from those in the '341 application in that the instant claims recite a method for reducing expression of a gene in a human cell the expression of which is inducible by an extracellular polypeptide that activates NF- κ B wherein the method comprises contacting the cell with a composition that diminishes NF- κ B activity whereas the claims in the '341 application recite a method for reducing expression of a gene in a human cell the expression of which is inducible by any external influence and wherein the method comprises within the cell inhibiting transmission of the signal so as to reduce expression of the gene. The instant claims are obvious however, because the '341 application specifically recites the species of extracellular polypeptides as external influences that activate NF- κ B and therefore said

extracellular polypeptides would have been an obvious species for activation of NF-KB. With regard to the instant claims reading on contacting the cell with the composition that diminishes NF-KB activity, it is noted that diminishing activity of NF-KB within the cell also results in inhibiting transmission of the signal (induced by NF-KB) in the cell since diminished NF-KB activity results in diminished signal transmission in the cell (as recited in the '341 claims).

This is a provisional obviousness-type double patenting rejection because the conflicting claims have not in fact been patented.

Applicants have responded to this rejection (see Applicants Remarks, filed 9 November 2009) by noting that the rejection is provisional and therefore defer discussion of the provisional rejection until the double patenting rejection is the only rejection remaining in the present application.

The rejection is therefore MAINTAINED.

Claim Rejections - 35 USC § 112 (1st paragraph)-maintained

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claim 89 STANDS rejected under 35 U.S.C. 112, first paragraph, as failing to comply with the written description requirement. The claim(s) contains subject matter which was not described in the specification in such a way as to reasonably convey to

one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention for reasons of record and presented herein.

Applicants response has been fully considered but is respectfully not found persuasive.

Initially it is noted that in the recent court decision *Ariad Pharms., Inc. v. Eli Lilly & Co.* Appeal from the US District Court for the District of MA (Decided 3 April 2009), Claims 80, 95, 144, and 145 (which depend from Claims 7-9 and 14) of U.S. Patent No. 6,410,516 (hereafter the '516 patent) were held to be unpatentable for failing to comply with the written description requirement. Because the pertinent claims of the '516 patent are very similar to the instant claim 89 (see ODP rejection above), and because the instant application 10/037,415 is a CONTINUATION of the '516 patent, the relevant criteria used for determining lack of written description for the '516 patent is compared to the instant claim 89 and applied below in a chart format for clarity of the record. The instant application fails to meet the criteria for written description set forth by the *Ariad* decision in at least the following ways as shown in the Table below:

Current Claim	Ariad Claims	Ariad v. Lilly decision factors/criterion	Reasoning as to why current claim does not meet criteria
89	80,95 144,145	Describes function but not what the composition is	The instant claim only defines the function.
		A vague functional description and an invitation for further research does not constitute written disclosure	The instantly claimed methods comprising the single step of reducing NF- κ B activity is not supported by written description because the specification of the '516 patent fails to adequately disclose how the claimed reduction of

		<p>NF-κB activity is achieved. The specification of the '516 patent as does the current application, hypothesizes three classes of molecules potentially capable of reducing NF-κB activity: specific inhibitors, dominantly interfering molecules, and decoy molecules. However, this disclosure amounts to little more than a research plan, and does not satisfy the patentee's quid pro quo as described in <u>Rochester</u>.</p> <p>In <u>Rochester</u>, very similar method claims were held invalid for lack of written description. <u>Id.</u> (holding patent invalid because "Rochester did not present any evidence that the ordinarily skilled artisan would be able to identify any compound based on [the specification's] vague functional description"); <u>see also Fiers v. Revel</u>, 984 F.2d 1164, 1170–71 (Fed. Cir. 1993) (holding a claim to a genus of DNA molecules not supported by written description of a method for obtaining the molecules); <u>cf. Eli Lilly</u>, 119 F.3d at 1567–68 (holding claims to a broad genus of genetic material invalid because the specification disclosed only one particular species). Ariad attempts to categorically distinguish <u>Rochester</u>, <u>Fiers</u>, and <u>Eli Lilly</u>, because in those</p>
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		<p>cases, the claims explicitly included the non-described compositions.</p> <p>Regardless of whether the claims recite a compound, the specification still must describe some way of performing the claimed methods. In the instant case, the specification suggests only the use of the three classes of molecules to achieve NF-κB reduction. Thus, to satisfy the written description requirement for the instant claims, the specification must demonstrate that Applicant possessed the claimed methods by sufficiently disclosing molecules capable of reducing NF-κB activity so as to "satisfy the inventor's obligation to disclose the technologic knowledge upon which the patent is based, and to demonstrate that the patentee was in possession of the invention that is claimed."<u>Capon</u>, 418 F.3d at 1357.</p>
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			specification does not adequately describe using those molecules to reduce NF- κ B activity.
		Because written description is determined as of the filing date-- April 21, 1989 in the Ariad case and in the instant case,--evidence of what one of ordinary skill in the art knew in 1990 or 1991 cannot provide substantial evidence to support adequate written description. <u>See Vas-Cath</u> , 935 F.2d at 1563–64 (holding that a written description analysis occurs “as of the filing date sought”).	Evidence of what one of ordinary skill in the art knew in 1990 or 1991 cannot provide substantial evidence to support adequate written description. <u>See Vas-Cath</u> , 935 F.2d at 1563–64 (holding that a written description analysis occurs “as of the filing date sought”).
		Predictability	Ariad explains that developing the subject matter of the '516 patent “required years of hard work, great skill, and extraordinary creativity—so much so that the inventors first needed to discover, give names to, and describe previously unknown cellular components as a necessary predicate for their inventions.” Lilly offered the undisputed expert testimony of David Latchman that the field of the invention was particularly unpredictable. Thus, this invention was made in a new and unpredictable field where the existing knowledge and prior art was scant. <u>See Capon</u> , 418 F.3d at 1359.

The description requirement of the patent statute requires a description of an invention, not an indication of a result that one might achieve if one made that invention. See *In re Wilder*, 736, F.2d 1516, 1521, 222 USPQ 369, 372-73 (Fed. Cir. 1984) (affirming rejection because the specification does “little more than outlin[e] goals appellants hope the claimed invention achieves and the problems the invention will hopefully ameliorate.”) Accordingly, it is deemed that the specification fails to provide adequate written description for the genus of the claims and does not reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the entire scope of the claimed invention.

Applicants response is to traverse the rejection. Applicants argue that the Examiner's support for the assertion of lack of written description that was based on the decision of the Court of Appeals of the Federal Circuit in *ARIAAD Pharmaceuticals, Inc et al v. Eli Lilly and Company*, 560 F.3d 1366 (Fed. Cir. 2009) and thus the rejection of the pending claims based on the support provided in the ARIAD decision has been rendered moot because Applicants submit that the Federal Circuit granted ARIAD's petition for rehearing *en banc* on August 21, 2009 and that therefore this petition decision “also vacated and rendered void the earlier ARIAD decision.

Applicants arguments have been fully considered but are not persuasive because Applicant has neither distinctly pointed out how Applicant believes the points made in the previous rejection are incorrect nor argued why the pending claim meets the requirement for written description under 35 U.S.C. 112, first paragraph, nor pointed out where the pending claims are described in the specification in such a way as to

reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention.

Conclusion

No claims are allowed.

THIS ACTION IS MADE FINAL. Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to CATHERINE HIBBERT whose telephone number is (571)270-3053. The examiner can normally be reached on M-F 8AM-5PM, EST.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Christopher Low can be reached on 571-272-0951. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Catherine Hibbert
Examiner AU1636

/ Christopher S. F. Low /
Supervisory Patent Examiner, Art Unit 1636